# PATENT COOPERATION TREATY

# **PCT**

# INTERNATIONAL PRELIMINARY EXAMINATION REPORT

REC'D	13	MAY	2005
WIPO			PCT

(PCT Artcle 36 and Rule 70)

Applicant's or agent's file reference		SeeNotification	nofTransmittalofInternational	Preliminary
PCA30425/PSC	FOR FURTHER ACTION	Examination R	eport (Form PCT/IPEA/416)	
International application No.	International filing date(day/mo	1	Priority date (day/month/yea	<b>)</b>
PCT/KR2003/000857	28 APRIL 2003 (28.04.2		09 JANUARY 2003 (09.01	.2003)
International Patent Classification (IPC IPC C07F 9/22, C07F Applicant				
POSTECH FOUNDATION	et al	orad by this Inter	national Preliminary Examini	ing Authority
and is transmitted to the applica	nt according to Article 36.			
2. This REPORT consists of a total of sheets, including this cover sheet.  This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).				
These annexes consist of a total of sheets.				
3. This report contains indications	s relating to the following items:			
I Basis of the repor	t			
II Priority				
III Non-establishmen	nt of opinion with regard to novelt	y, inventive step	and industrial applicability	
IV Lack of unity of i	nvention			
Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement				
VI Certain documen	VI Certain documents cited			
VII Certain defects in	the international application			•
VIII Certain observations on the international application				
Date of submission of the demand	Da	te of completion o	of this report	
26 JULY 2004 (	26.07.2004)	14 APRIL 2	2005 (14.04.2005)	
Name and mailing address of the IPI	EA/KR Au	nthorized officer		FILE
Korean Intellectual Property Office 920 Dunsan-dong, Seo-gu, Daejeon 302-701, Republic of Korea		MOON, Sun I	Ieup	
Facsimile No. 82-42-472-7140	Te	elephone No. 82-	42-481-5543	

# INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International aplication No. PCT/KR2003/000857

I.	Bas	is of th	e report	
1.	Wit	h regar	to the elements of the international application:*	
		] the in	ternational application as originally filed	
	$\boxtimes$	4	escription:	
			s 1-25 s NONE , filed with the demand	
			s NONE , filed with the letter of	
	K	_	claims:	
	$\boxtimes$		, as originally filed	
		page	NONE , as amended (together with any statment) under Article 19	
			NONE, the with the letter of 08 MARCH 2005	
	<u></u>	 m		
	$\succeq$	the	drawings: , as originally filed	
		page	NONE , filed with the definant	
		page	NONE , filed with the letter of	
	$\geq$		sequence listing part of the description:	
Ì		page	S 1-0	
		pag	NONE , filed with the letter of	
<b>]</b> .	٠			
2	. V	Vith reg	ard to the language, all the elements marked above were available or furnished to this Authority in the language in which	
	th	ne inter	national application was filed, unless otherwise indicated under this item.  ements were available or furnished to this Authority in the following language <u>English</u> which is	
	Т			
	L		e language of a translation furnished for the purposes of international search (under Rule 23.1(b)).	
1	$\triangleright$	≤ th	e language of publication of the international application(under Rule 48.3(b)).	
			e language of the translation furnished for the purposes of international preliminary examination (under Rules 55.2 and/	
1			55.3).	
	3.	With r	egard to any nucleotide and/or amino acid sequence disclosed in the international application, the international nary examination was carried out on the basis of the sequence listing:	
	D	_	ontained inthe international application in written form.	
	5	_	led together with the international application in computer readable form.	
1	Г		urnished subsequently to this Authority in written form.	
l	L [	_	urnished subsequently to this Authority in computer readable form	
l	The second that the subsequently formished written sequence listing does not go beyond the disc losure in the			
	international applications as filed has been furinshed.			
	ľ	$\nabla$	he statement that the information recorded in computer readable form is identical to the written sequence using has	
	٤		een furnished.	
	4 F	√ 1	he amendments have resulted in the cancellation of:	
Ţ,	4. [	_ لاحث	<del>- 7</del> 1	
		<u> </u>	the description, pages NONE the claims, Nos. NONE	
-		Į.		
	_		the drawings, sheet NONE	
	5. 	□ <sup>,</sup>	This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box(Rule 70.2(c)).**	
	1	Replace in this c and 70.	ement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to opinion as "originally filed." and are not annexed to this report since they do not contain amendments (Rules 70.16 17).	
	**	Any rej	placement sheet containing such amendments must be referred to under item I and annexed to this report.	

#### INTERNATIONAL PRELIMINARY EXAMINATION

International aplication No.
PCT/KR2003/000857

V. Reasoned statement under Article 35(2) with regard to	novelty, inventive step or industrial applicability;
citations and explanations supporting such statement	

1.	Statement			
	Novelty (N)	Claims	1 - 2	YES
	1.0.0.0, (1.7)	Claims	NONE	NO.
	Inventive step (IS)	Claims	1 - 2	YES
İ	mvomvo stop (15)	Claims	NONE	NO
	Industrial applicability (IA)	Claims	1 - 2	YES
	••	Claims	NONE	

- 2. Citations and explanations (Rule 70.7)
  - 1. Reference is made to the following documents:
  - D1: US 5969128 A(La Region Wallone) 19 OCTOBER 1999
  - D2: Marie-Laurence Fontanel et al. 'Sterical recongition by T4 polyneucl eotide kinase of non-nucleosidic moieties 5'-attached to oligonucleotides', Nucleic Acids Research, Oxford Univ. Press, 1994, v.22, no.11, pp.2022-7
  - D3: De Vos, M.J. et al. 'New ono-nucleosidic phosphoamidites for the solid-phase multilabeling of oligodeoxyribonucleotides', Nucleosides&nuclotides, 1994, v.13, no.10, pp.2245-65
  - D4: Wilk, Andrzej et al. 'Backbone-modified oligonucleotides ocntain ing a butanediol-1,3 moiety as a vicarious segment or the deoxyribosyl moiety-synthesis and enzyme studies', Nucleic Acids Research, Oxford Univ. Press, 1990, v.18, no.18, pp.2065-8
  - 2. Novelty and Inventive Step

The subject matters of the present invention relate to a novel phoshoramidite compound which can be used in the synthesis of DNA variants by introducing a desired functional group, e.g, benzyl glycolate, lithocholic acid, pentaerythritol, and dendrimer.

The prior art documents, D1-D4 relate to the phosphoramidite compound of 1,3-butanediol functional substitutent.

None of the prior art documents disclose or teach various functional groups like benzyl glycolate, lithocholic acid, pentaerythritol, and dendrimer. In the light of the chemical structural difference among the present invention and D1-D4, it is not obvious to the skiled in the art to select the functional groups of the present invention which essentially differ from those of D1-D4.

Therefore, the subject matter of claims 1-2 is considered to be novel and to involve an inventive step under PCT Article 33(2)-(3).

3. Industrial Applicability

The subject matter of the claims 1-2 is considered to be industrially applicable under PCT Article 33(4).

# Amendment to the Claims

Please replace originally filed pages 26 and 27 with the enclosed new pages 26 and 27, wherein claims 1 and 2 have been amended.

AMENDED SHEET (ART. 34)

# What is claimed is:

1. A phosphoramidite compound of formula (I), (II), (III), (IV) or

(II)

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AMENDED SHEET (ART. 34)

wherein;

R is a dimethoxytrityl (DMTr), levulinyl (Lev) or tert-butyldimethylsilyl (TBDMS) group.

5 2. The compound of claim 1, which is selected from the group consisting of:

O-((2-cyanoethyl)-N,N-diisopropyl-phosphoramidite)-benzylglycolate;

O-DMTr-((2-cyanoethyl)-N,N-diisopropyl-phosphoramidite)-

10 lithocholic alcohol;

O-tri-DMTr-((2-cyanoethyl)-N,N-diisopropyl-phosphoramidite)-pentaerithritol;

O-DMTr-O-di-Lev-O-((2-cyanoethyl)-N,N-diisopropyl-phosphoramidite)-pentaerithritol; and

O-DMTr-O-Lev-O-TBDMS-((2-cyanoethyl)-N,N-diisopropyl-phosphoramidite)-pentaerithritol.